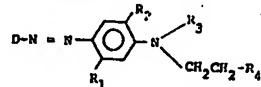


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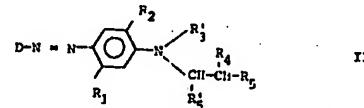
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(54) Improvements in or relating to
organic compounds

(57) The present invention relates to a
mixture of disperse dyestuffs contain-
ing at least one compound of formula I,



and at least one compound of formula II



in which D, is a disperse dye diazo
component radical,

R₁, is hydrogen, alkyl, alkoxy or acyla-
mino,

R₂, is hydrogen, alkyl or alkoxy which
is optionally substituted,

R₃, is hydrogen, alkenyl, chloro- or
bromo-alkenyl, linear alkyl which is
optionally substituted,

R₄, is hydrogen, alkenyl, chloro- or
bromo alkenyl, linear or branched alkyl,
which is optionally substituted,

R₄, is a sulphur-free acyloxy or
acyloxy-carbonyl radical,
one of R₅ and R₆ is hydrogen and the
other is methyl,

the D's R₁'s, R₂'s and R₄'s being
identical, which mixture contains 15 to
85% compound of formula I and 85 to
15% compound of formula II, and has
good dispersion stability and is useful
for dyeing disperse dye dyeable sub-
strats with good build-up power and
gives dyeings have good wet-fastness.

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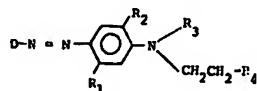
SPECIFICATION

Improvements in or relating to organic compounds

5 The present invention relates to mixtures of disperse dyestuffs which mixtures possess improved properties over the individual dyestuffs.

More particularly, the present invention provides a mixture of disperse dyestuffs containing at least one compound of formula I,

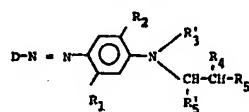
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I

15 and at least one compound of formula II

15



II

20

20

in which D, is a disperse dye diazo component radical,

R₁, is hydrogen, alkyl, alkoxy or acylamino,

R₂, is hydrogen, alkyl or alkoxy which is optionally mono-substituted by a substituent selected from halogen, cyano, acyl, acyloxy, hydroxyl, alkoxy, phenyl and phenoxy,

25 R₃, is hydrogen, alkenyl, chloro- or bromo-alkenyl, linear alkyl which is optionally substituted by up to two substituents selected from halogen, acyloxy, hydroxyl, and alkoxy or mono-substituted by a cyano, acyl, phenyl or phenoxy group,

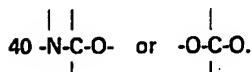
R₄, is hydrogen, alkenyl, chloro- or bromo-alkenyl, linear or branched alkyl, which is optionally substituted by up to two substituents selected from halogen, acyloxy, hydroxyl, and alkoxy or mono-substituted by

30 a cyano, acyl, phenyl or phenoxy group,

R₅, is a sulphur-free acyloxy or alkoxy-carbonyl radical, one of R₅ is R₆ is hydrogen and the other is methyl, the D's, R₁'s, R₂'s and R₄'s being identical, any alkyl

35 groups or moieties and any alkoxy groups in such dyestuffs contain 1 to 8 carbon atoms, any alkenyl groups 2 to 4 carbon atoms, and said mixture contains 15 to 85% compound of formula I and 85 to 15% compound of formula II.

It will be appreciated that the compounds of formula I and formula II are free from acetal groupings i.e.



40

By halogen is meant chlorine, bromine or iodine.

The preferred halogens are chlorine and bromine.

45 By 'acyl' is meant groups of formula R-Y- and R'-Z wherein R is a alkyl, alkenyl, C₅₋₇ cycloalkyl, phenyl or a heterocyclic group, (preferably alkyl, alkenyl (C₅₋₇) cycloalkyl or phenyl), which alkyl is optionally mono-substituted by halogen, hydroxy, (C₁₋₂) alkoxy, phenyl or phenoxy and which phenyl is optionally substituted by up to two substituents selected from halogen, (C₁₋₂) alkyl and (C₁₋₂) alkoxy,

Y is -O-CO-, or -OSO₂(-O-bound to R) or -SO₂-

50 R' is hydrogen or has one of the significances of R, Z is -CO-, -NR'CO- or -NR'SO₂(-N bound to R'), and R'' is hydrogen or has one of the significance of R

and acyloxy and acylamino are to be understood accordingly.

When the alkyl and/or alkoxy groups in the dyestuffs are substituted they are preferably monosubstituted or are substituted by a hydroxyl group and a further substituent. Preferably the alkyl and alkoxy radicals contain 1 to 4, more preferably 1 or 2 carbon atoms. Preferred alkenyl groups contain 3 carbon atoms. The preferred cycloalkyl is cyclohexyl.

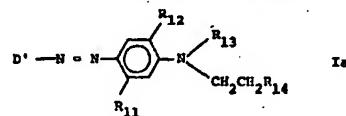
Preferred diazo component radicals are those of the benzene series (optionally disazoc component radicals) and diazo component radicals of the benzothiazolyl, thiophene or thiazolyl series.

60 More preferably, D is thiophene-2 substituted by up to two nitr groups; benzothiazolyl-2 in which the benzene nucleus is substituted by up to two substituents selected from chlorine and nitro; or phenyl substituted in the para position by nitro, alkylsulphonyl, aminosulphonyl, mono- or dialkylaminosulphonyl or phenylazo (in which the benzene nucleus is optionally substituted by up to three substituents selected from methyl, methoxy, chlorine, bromine and nitro) and optionally bearing up to two further substituents selected from halogen, nitro, cyano or alkylsulphonyl.

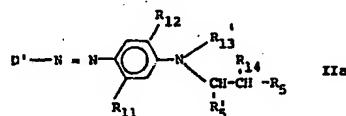
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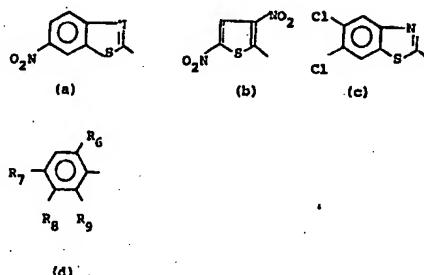
Representative mixtures are those containing at least one compound formula i.e.



and at least one compound of formula IIa



15 In which D' is a radical of formula (a), (b), (c) or (d)



	wherein R ₆	Is hydrogen, chlorine, bromine, iodine, nitro or cyano,
30	R ₇	is nitro, (C ₁₋₂)alkylsulphonyl, amino-sulphonyl, mono- or di(C ₁₋₂) alkylamino-sulphonyl or phenylazo in which the benzene ring is optionally substituted by up to three substituents selected from methyl, methoxy, chlorine, bromine and nitro,
	R ₈	is hydrogen, chlorine or bromine,
35	R ₉	is hydrogen, chlorine, bromine, iodine, cyano or (C ₁₋₂) alkylsulphonyl, with the proviso that one of R ₈ and R ₉ is hydrogen,
	R ₁₁	is hydrogen, methyl, methoxy, ethoxy, formylamino, alkyl (C ₁₋₄) carbonylamino, alkyl (C ₁₋₂) carbonylamino monosubstituted by hydroxyl, chlorine, bromine, cyano, phenyl, (C ₁₋₃) alkoxy or phenoxy, alkoxy-(C ₁₋₄) carbonylamino, ethoxycarbonylamino monosubstituted by hydroxyl, (C ₁₋₂ d ₂₋₃) alkoxy or (C ₁₋₂) alkoxyethoxy, benzoylamino or (C ₁₋₂) alkylsulphonylamino,
40	R ₁₂	is hydrogen, methyl, (C ₁₋₂) alkoxy, (C ₁₋₂)-alkoxyethoxy or (C ₁₋₂) alkoxy monosubstituted by chlorine, bromine, cyano or phenyl,
	R ₁₃	is hydrogen, linear (C ₁₋₄) alkyl, (C ₂₋₃)-alkenyl, linear (C ₂₋₃) alkyl mono-substituted by (C ₁₋₂) alkoxy, cyano, formyloxy, alkyl (C ₁₋₄) carbonyloxy, alkoxy (C ₁₋₃) carboxyl, benzoyloxy or alkoxy (C ₁₋₃) carbonyloxy,
45	R' ₁₃	has one of the significances of R ₁₃ but where the alkyl and substituted alkyl radicals are linear or branched,
	R ₁₄	is formyloxy, alkyl (C ₁₋₄) carbonyloxy, benzoyloxy, phenoxyacetoxyl, alkoxy (C ₁₋₄)-carbonyloxy, alkoxy (C ₁₋₄) carbonyl, alkoxy (C ₁₋₂) ethoxycarbonyl, alkoxy (C ₁₋₂) ethoxycarbonyloxy, alkoxy (C ₁₋₂) alkyl (C ₁₋₂) carbonyloxy, mono- or di (C ₁₋₄) alkyl) aminocarbonyloxy or phenylaminocarbonyloxy wherein the benzene nucleus is optionally substituted by up to two substituents selected from chlorine, bromine, methyl and methoxy,
50		

whereby in such mixture D', R₁₁, R₁₂, and R₁₄ in the compound of formula Ia is identical with that of the compound of formula IIa.

More preferred mixtures are those wherein D' is a group of formula (d), especially those wherein R₆ is R'₆, where R'₆ is hydrogen, chlorine, bromine or nitro, R₇ is nitro, R₈ is hydrogen and R₉ is R'₉ where R'₉ is hydrogen, chlorine, bromine, cyano or methylsulphonyl.

Even more preferred mixtures are those wherein D' is a group $\text{R}_8\text{R}_9\text{R}_{10}$ wherein R_8 is R'_8 , R_9 is nitro R_8 is hydrogen, R_9 is R'_9 , R_{10} is R'_{10} where R'_9 is hydrogen, methyl, alkyl (C_{1-2}) carbonylamin, chloro- or bromoalkyl- (C_{1-2}) carbonylamin, alkoxy (C_{1-2}) carbonylamin, alkyl xy-(C_{1-2}) eth ycarbonylamino, R_{12} is R'_{12} , where R'_{12} is hydrogen, methoxy or eth oxy, R_{13} is R'_{13} , where R'_{13} is linear (C_{2-3}) alkyl, allyl or linear (C_{2-3}) alkyl monosubstituted by cyan, alkyl (C_{1-2}) carbonyloxy, alkyl xy- (C_{1-2})- carbonyl or alkoxy (C_{1-2}) carbonyl xy, R'_{13} has one of the significances of R_{13a} but where the partially substituted (C_{2-3}) alkyl may be linear or branched and R_{14} is alkyl- (C_{1-2}) carbonyloxy or alkoxy (C_{1-2}) carbonyl xy.

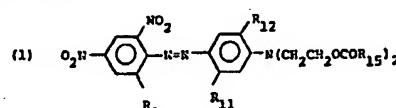
The preferred coupling component in the mixtures of the invention are those where R₆ is hydrogen, R₅ is methyl, R₁ is R₁₁, R₂ is R₁₂, R₃ is R₁₃, R₄ is acyl xy specially alkyl (C₁₋₂) carbonyloxy or

alkoxy-(C₁₋₂) carbonyloxy, especially those wherein R₁₁ is R'₁₁, R₁₂ is R'₁₂, R₁₃ is R'_{13a} and R'₁₃ has one or two significances R'_{13a} but where the optionally substituted alkyl may be linear or branched.

The following mixtures of the compounds of formula Ia and IIa are most preferred

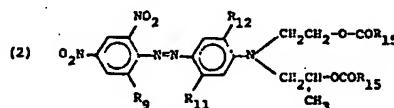
- (i) those where D' is a radical (d) wherein R₆ is chlorine or bromine, especially chlorine, R₇ is nitro, R₈, R₉, R₁₁ and R₁₂ are hydrogen, R₁₃ and R'₁₃ are cyanoethyl, R₁₄ is alkyl (C₁₋₂) carbonyloxy, especially acetoxy, R₅ is methyl and R'₅ is hydrogen; 5
- (ii) those where D' is a radical (d) wherein R₆, R₈, R₁₁ and R₁₂ are hydrogen, R₇ is nitro, R₉ is cyano, R₁₃ and R'₁₃ are cyanoethyl, R₁₄ is alkyl (C₁₋₂) carbonyloxy, especially acetoxy, R₅ is methyl and R'₅ is hydrogen; 10
- (iii) those wherein D' is a radical (d) wherein R₆ and R₉, independently, are chlorine, or bromine, especially chlorine, R₇ is nitro, R₈, R₁₁ and R₁₂ are hydrogen, R₁₃ and R'₁₃ are cyanoethyl, R₁₄ is alkyl (C₁₋₂) carbonyloxy, especially acetoxy, R₅ is methyl and R'₅ is hydrogen; 10
- (iv) a mixture of the following three dyes:

15



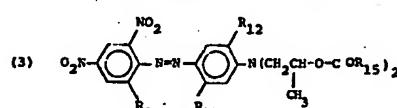
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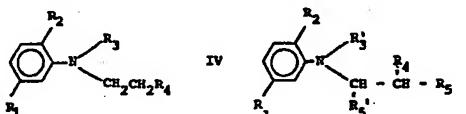


25

In which each R₉ is chlorine, bromine or iodine, especially bromine, each R₁₁ is alkyl (C₁₋₂) carbonylamino, especially acetylarnino, each R₁₂ is (C₁₋₂) alkoxy, especially ethoxy and each R₁₅ is (C₁₋₂) alkyl, especially methyl, especially a mixture containing approximately 25% (1) 50% (2) and 25% (3).

The preferred method of preparing the mixtures according to the invention is to couple a diazotised amine with a mixture containing at least one compound of formula IV and at least one compound of formula V,

35



35

40 whereby at least 15 mol % of a compound of formula IV and at least 15 mol % of a compound of formula V is present. 40

Diazotization and coupling are carried out in accordance with known methods.

It will also be appreciated that the mixtures of the invention may also be obtained by admixing dyes of formula I and II together, preferably at the time of making the dyeing preparation.

45 The dyestuff mixtures according to the present invention are made up into dyeing preparations by known methods, for example by grinding in the presence of disperse agents and/or fillers, followed by drying. It will be appreciated that when the mixtures are obtained by admixing dyes of formula I and II, the dyeing preparation can be made by admixing the already ground dyes of formula I and II or by grinding the already mixed dyes of formula I and II.

50 The dyestuff preparations containing the mixtures according to the present invention are useful for dyeing or printing textiles comprising or consisting of synthetic, semi-synthetic, hydrophobic, high-molecular weight organic substrates from aqueous suspensions. Preferred substrates are textiles of linear, aromatic polyesters, cellulose 2/2 acetate, cellulose triacetate and synthetic polyamides. Dyeing, padding or printing may be carried out in accordance with known methods, for example as described in French Patent No. 50

55 1,445,371.

It is known that some disperse dyes exist in the thermoinstable form which causes problems in that they do not build-up in the fibre which leads to uneven dyeings having poor rubbing and wash fastness and which causes special difficulties for cross-wound spool dyeing. Such dyestuffs can be converted to the therm stable form by heat-treatment. However with the dyestuff preparations made with some of the mixtures according to the present invention the above-mentioned disadvantages do not occur and thus, the further conversion e.g. heat-treatment step is unnecessary. 60

Further, the mixtures of the invention have good dispersion stability, have good build-up power and give dyeings with good rubbing and fastnesses.

The following Examples further serve to illustrate the invention. In the Examples all parts are by weight, 65 unless otherwise stated, and all temperatures are in degree centigrade.

55

65

Example 1

17.3 Parts 2-chloro-4-nitroanilin are stirred overnight in a mixture of 44 parts conc. hydrochloric acid and 100 parts water. 60 Parts ice are added thereto and a solution of 6.9 sodium nitrite in 25 parts water are added dropwise over a period of 30 minutes at 0 - 3°. After 2½ hours when diazotization is completed some undissolved product is filtered off and a mixture of 11.6 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)-aminobenzene, 12.3 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)-aminobenzene and 20 parts acetic acid are added dropwise over a period of 45 minutes at 0 to 3° to the clear diazonium salt solution. The coupling reaction is completed by the addition of 50 parts sodium acetate. The precipitated dyestuff mixture is filtered, washed with 4000 parts water and dried.

10 The dyestuff mixture without conversion to the thermostable form may be up into a dyeing preparation in accordance with known methods with the addition of a dispersing agent, e.g. a commercially available ligninsulphonate which preparation has good dispersion stability.

Example 2

15 16.3 parts 2-cyano-4-nitroaniline are added at 5 to 10° to 100 parts (95%) sulphuric acid. At the same temperature, 32 parts (40%) nitrosylsulphuric are added dropwise and the whole is stirred for 3 hours. The so obtained diazonium salt solution is poured onto a mixture of 300 parts ice and 100 parts water. Some undissolved product is filtered off and a mixture of 11.6 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)-aminobenzene, 12.3 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)-aminobenzene and 30 parts acetic acid is added dropwise at 0 to 3° to the clear solution. After 3 hours the precipitated dyestuff mixture is filtered, washed free of acid and dried. Dyestuff preparations made therefrom without conversion have good dispersion stability.

Example 3

20 25 A mixture of 16.2 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)aminobenzene, 7.4 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)aminobenzene and 30 parts acetic acid are added dropwise to the clear diazonium salt solution obtained as described in Example 2 at 0 to 3°. The precipitated dyestuff mixture is filtered and washed free from acid. The presscake is subsequently stirred in 300 parts water. The homogenous suspension is put in a steel autoclave and is stirred for 3 hours at 105 - 110°. After cooling and filtering, a 30 thermostable dyestuff mixture is obtained which may be made up into useful dyestuff preparations.

Example 4

35 35 13.8 Parts 4-nitroaniline are dissolved in 80 parts water at 70 - 80° with the addition of 28 parts conc. hydrochloric acid. The solution is cooled to 0 to 2° by adding 200 parts ice and diazotization is quickly carried out with a solution of 6.9 parts sodium nitrite in 14 parts water. Some solid product is filtered off and over a period of 60 minutes a mixture of 13.9 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)aminobenzene, 9.85 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)aminobenzene and 20 parts acetic acid is added dropwise at 0 to 3° to the clear diazonium salt solution. After the addition of 40 parts sodium acetate coupling is complete and the precipitated dyestuff mixture is filtered, washed and dried. The mixture, without conversion, is made into a 40 finely divided dyeing preparation with the addition of dispersing agent, which preparation gives orange dyeings on polyester fibres.

Example 5

45 45 At 0 to 3° a mixture of 18.6 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)aminobenzene, 4.9 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)aminobenzene and 20 parts acetic acid are added dropwise over a period of 45 minutes to the diazonium-salt solution prepared as described in Example 1. 50 Parts sodium acetate are added to complete the coupling reaction. The precipitated dye mixture is filtered, washed with 4000 parts water and dried in vacuo for at least 48 hours at 100-105°.

50 After heat treatment to convert the mixture to the thermostable form the dye mixture is made up into an excellent finely dispersed dyestuff preparation.

Example 6

55 55 69 parts sodium nitrite are added portionwise to 1200 parts conc. sulphuric acid at 60-70° with energetic stirring. Stirring is continued for a further 10 minutes and the mixture is cooled to 30° whereupon 207 parts 2,6-dichloro-4-nitroaniline are added. Diazotization is complete after 3 hours. The yellow-brown solution is poured onto a mixture of 3400 ice and 2500 parts water. A mixture of 116 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)aminobenzene, 123 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)aminobenzene and 200 parts acetic acid are added dropwise with stirring at 0 to 5° to the obtained aqueous diazonium-salt solution. The precipitated dyestuff mixture is filtered, washed acid-free, dried and, without conversion, the therm stable forms is made up in accordance with known methods, with the addition of a ligninsulphonate into a fine dispersion. The preparation has very good dispersion stability combined with good build-up power on polyester material.

60 65 A mixture of 46 parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)aminobenzene, 197 parts N-(2'-acetoxypropyl)-

N-(2'-cyan ethyl)aminobenzene and 200 parts acetic acid are dropped into the diazonium salt solution of Example 6 at 0 to 5°. The precipitated dye is filtered and washed acid free. A dispersion stable dye mixture is obtained which gives satisfactory dyeings without conversion to the thermostable form.

5 Example 8

600 Parts water and 194 parts 1-acetyl amino-4-ethoxy-3-amino benzene are put in an autoclave which is rendered oxygen free by passing nitrogen therethrough. The whole is heated to 55° and the amine is reacted for 2 hours with a mixture consisting of 52.8 parts ethyleneoxide and 69.6 parts propylene oxide whereupon the temperature rises to 62°. The mixture is stirred for 6 hours at 60° and then cooled to 5°. Excess ethylene-
10 and propyleneoxide are removed by means of a water jet pump and then passing nitrogen therethrough. The autoclave is opened and the crystalline precipitated product is filtered, washed with a little cold water and dried in vacuo at 60°.

148 Parts of the dried product are dissolved in 150 parts glacial acetic acid. At 80-90° over a period of 1 hour 122 parts acetic acid anhydride are added dropwise. The mixture is then heated to 110° and stirred for 3
15 hours at this temperature until there are no more traces of nonesterified material (tested by thin layer chromatography). The mixture of coupling components is cooled to room temperature.

With energetic stirring 31.1 parts sodium nitrite are added at 60 to 70 to 540 parts conc. sulphuric acid.
Stirring is continued for a further 10 minutes at 60°, the mixture cooled to 10° and 117.9 parts
2-bromo-4,6-dinitroaniline are added thereto. Diazotization is complete after 3 hours. The diazonium solution
20 is added to a mixture of all the coupling component obtained above, 450 parts glacial acetic acid, 3000 parts ice and 9 parts aminosulphonic acid at -5 to +2° over a period of 30 to 40 minutes. The dyestuff precipitates quickly and the solution is stirred at 0 to 2°C until no further diazo compound is detectable. The dye suspension is adjusted to pH 6 to 7 by the addition of 30% sodium hydroxide solution. Then the solution is heated to 85° over a period of an hour and is kept at this temperature for an hour followed by heating to 90°.
25 and stirred for a further hour whereby the dyestuff components convert to the thermostable form. The dye is then filtered, washed with 5000 parts hot water and dried in vacuo at 80°.

D Preparation of Dispersion

In a flask, 78 parts of the dye mixture of Example 8 and 222 parts commercially available ligninsulphonate
30 dispersing agent in 100 to 150 parts water are ground with 800 parts by volume silica quartz beads for 6 hours (2000-2500 R.p.m.). The whole is diluted with 300 parts water and suction filtered. Any dyestuff and/or dispersing agent remaining on the beads is rinsed into the flask with a little water. The dispersion is adjusted to pH 6 to 6.5 with phosphoric acid and spray-dried (hot air 130°).

35 Example 9

Preparation of Dye A

232 Parts N-(2'-acetoxyethyl)-N-(2'-cyanoethyl)-aminobenzene and 200 parts acetic acid were poured into the diazonium salt solution produced as described in Example 6 at 0 to 5°. The precipitated dye is filtered
40 washed free of acid and dried.

Preparation of Dye B

The diazonium salt solution of Example 6 is coupled with 246 parts N-(2'-acetoxypropyl)-N-(2'-cyanoethyl)aminobenzene in 200 parts acetic acid. The dye is filtered, washed and dried.

45 Preparation of Dyeing Formulation

22.2 Parts dye A and 70.8 parts dye B are ground and dried with 207 ligninsulphonate as described above under D. The powdered product obtained has notable dispersion stability.

The mixtures given in the following Table may be prepared in accordance with the procedures of the foregoing Examples.

TABLE

Ex. No.	General Formula	1st Component Mol %	2nd Component Mol %	3rd Component Mol %	shade on Polyester
10		X = H 50	X = CH ₃ 50	-	navy-blue
11		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	brown
12		X = H 40	X = CH ₃ 60	-	blue
13		X = H 50	X = CH ₃ 50	-	orange
14		X = H 70	X = CH ₃ 30	-	yellow-brown
15		X = H 55	X = CH ₃ 45	-	navy-blue
16		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = CH ₃ ca. 50 X ₂ = H	scarlet
17		X = H 50	X = CH ₃ 50	-	orange
18		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	yellow-brown
19		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	yellow-brown
20		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	navy-blue
21		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	red
22		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	scarlet
23		X ₁ = H ca. 25 X ₂ = H	X ₁ = CH ₃ ca. 25 X ₂ = CH ₃	X ₁ = H ca. 50 X ₂ = CH ₃	red

Ex. No.	General Formula	1st Component : Mol %	2nd Component : Mol %	3rd Component : Mol %	shade on Polyester
24		X = H 45	X = CH ₃ 55	-	red
25		X = H 50	X = CH ₃ 50	-	red
26		X = H 50	X = CH ₃ 50	-	ruby
27		X = H 30	X = CH ₃ 70	-	red
28		X = H 35	X = CH ₃ 45	-	red
29		X = H 50	X = CH ₃ 50	-	reddish violet
30		X ₁ = H ca.25 X ₂ = H ca.25	X ₁ = CH ₃ ca.25 X ₂ = CH ₃ ca.50	X ₁ = H ca.50 X ₂ = CH ₃	blue
31		X = H 50	X = CH ₃ 50	-	scarlet
32		X ₁ = H ca.25 X ₂ = H ca.25	X ₁ = CH ₃ ca.25 X ₂ = CH ₃ ca.50	X ₁ = H ca.50 X ₂ = CH ₃	red
33		X ₁ = H ca.25 X ₂ = H ca.25	X ₁ = CH ₃ ca.25 X ₂ = CH ₃ ca.50	X ₁ = H ca.50 X ₂ = CH ₃	yellow- brown
34		X = H 50	X = CH ₃ 50	-	red
35		X ₁ = H ca.25 X ₂ = H ca.25	X ₁ = CH ₃ ca.25 X ₂ = CH ₃ ca.50	X ₁ = H ca.50 X ₂ = CH ₃	ruby
36		X = H 50	X = CH ₃ 50	-	reddish navy- blue
37		X ₁ = H ca.25 X ₂ = H ca.25	X ₁ = CH ₃ ca.25 X ₂ = CH ₃ ca.50	X ₁ = H ca.50 X ₂ = CH ₃	scarlet
38		X = H 70	X = CH ₃ 30	-	red

Ex. No.	General Formula	1st Component : Mol %	2nd Component : Mol %	3rd Component : Mol %	state on Polyester
39		x = H 55	x = CH3 45	-	orange
40		x = H 60	x = CH3 40	-	brown
41		x = H 50	x = CH3 50	-	red
42		x = H 25	x = CH3 75	-	scarlet
43		x = H 50	x = CH3 50	-	scarlet
44		x1 = H ca. 25	x1 = CH3 ca. 25	x1 = H ca. 50	navy-blue
45		x1 = H ca. 25	x1 = CH3 ca. 25	x1 = H ca. 50	brown
46		x = H 30	x = CH3 70	-	red
47		x = H 85	x = -CH3 15	-	brown
48		x = H 20	x = CH3 80	-	scarlet
49		x = H 65	x = CH3 35	-	yellow-brown
50		x = H 35	x = CH3 65	-	navy-blue
51		x = H 50	x = CH3 50	-	navy-blue
52		x = H 15	x = CH3 85	-	yellow-brown
53		x = H 80	x = CH3 20	-	ruby

Ex. No.	General Formula	1st Component Mol %	2nd Component Mol %	3rd Component Mol %	shade of Polyester
5	54	X = H 60	X = CH3 40	-	orange
10	55	X = H 50	X = CH3 50	-	navy-blue
15	56	X1 = H ca. 25 X2 = H	X1 = CH3 ca. 25 X2 = CH3	X1 = H ca. 50 X2 = CH3	ruby
20	57	X = H 75	X = CH3 25	-	scarlet

20

Application Example A
Dyeing Preparation II

93 Parts of Dyestuff A above are ground with 207 parts ligninsulphonate as described above under D.

25 Dyeing Preparation III

93 Parts dyestuff B above are ground with 207 parts ligninsulphonate as described above under D.

1000 Parts polyester material are pre-wetted for 5 minutes at 60° with 30 parts ammonium sulphate dissolved in 14 litres water and adjusted to pH5 with 85% formic acid. 31.6 Parts of dyeing preparation II and 8.4 parts dyeing preparation III dispersed in 1 litre water are added thereto. The dyebath is heated over 30 minutes to 130° and dyeing is conducted for 60 minutes. The dyebath is cooled to 80°, the dyed substrate removed, rinsed and dried. An even, rubbing-fast yellow-brown dyeing is obtained.

Application Example B

3 Parts of the preparation D above are stirred in 4000 parts water at 60°. With this dyebath 100 parts polyester fibre is dyed with the addition of 20 parts ortho-phenylphenol for 1 hour at 98°. After cooling, rinsing, soaping and drying a navy-blue dyeing with good fastnesses is obtained.

Application Example C

7 Parts of the mixture of Example 1 are ground to a fine powder with 4 parts sodiumdinaphthylmethanesulphonate, 4 parts sodiumcetyl sulphate and 5 parts anhydrous sodium sulphate in a ball mill for 48 hours.

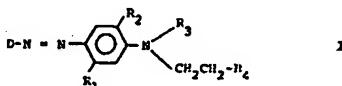
40 1 Part of this dyeing preparation is pasted with a little water and the suspension is added to a dyebath containing 3 parts sodiumlaurylsulphate in 4000 parts water. 100 Parts scoured polyester (liquor to goods ratio 40:1) are added to the bath at 40 to 50°, 20 parts of a chlorobenzene are added, the bath is slowly heated to 100° and dyeing is carried out for 1 to 2 hours at 95-100°. The even red dyeing which is washed, soaped, washed again and dried has good fastnesses.

45 In analogy to the above procedures the dyestuff mixtures of Examples 2 to 57 may be used to dye polyester fibres.

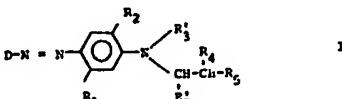
CLAIMS

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1. A mixture of disperse dyestuffs containing at least one compound of formula I,



and at least one compound of formula II



in which

D, is a disperse dy diazo component radical,
65 R1, is hydrogen, alkyl, alkyloxy or acylamin,

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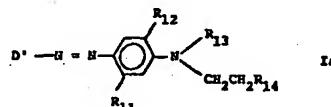
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- R₂ is hydrogen, alkyl or alkoxy which is optionally mono-substituted by a substituent selected from halogen, cyan, acyl, acyl xy, hydroxyl, alkoxy, phenyl and phenoxy,
R₃ is hydrogen, alkenyl, chloro- or bromo-alkenyl, linear alkyl which is optionally substituted by up to two substituents selected from halogen, acyloxy, hydroxyl, and alkoxy or mono-substituted by a cyano, acyl, phenyl or phenoxy group,
5 R₃' is hydrogen, alkenyl, chloro- or bromo alkenyl, linear or branched alkyl, which is optionally substituted by up to two substituents selected from halogen, acyloxy, hydroxyl, and alkoxy or mono-substituted by a cyano, acyl, phenyl or phenoxy group,
R₄ is a sulphur-free acyloxy or alkoxy carbonyl radical,
10 one of R₅ or R₆ is hydrogen and the other is methyl, the D's R₁'s R₂'s and R₄'s being identical, any alkyl groups or moieties and any alkoxy groups in such dyestuffs contain 1 to 8 carbon atoms, any alkenyl groups 2 to 4 carbon atoms, and said mixture contains 15 to 85% compound of formula I and 85 to 15% compound of formula II.
2. A mixture according to Claim 1, in which D is a radical of the benzene series, benzothiazolyl, thiazolyl
15 or thiophene series.
3. A mixture according to Claim 1 or Claim 2, in which D is thiophene-2-substituted by up to two nitro groups; benzothiazolyl-2 in which the benzene nucleus is substituted by up to two substituents selected from chlorine and nitro; or phenyl substituted in the para-position by nitro, alkylsulphonyl, aminosulphonyl, mono- or dialkyl-amino sulphonyl or phenylazo (which phenylazo is optionally substituted by up to three
20 substituents selected from methyl, methoxy, chlorine, bromine and nitro) and optionally bearing up to two further substituents selected from halogen, nitro, cyano or alkylsulphonyl, any alkyl and alkoxy groups in said dyestuffs contain 1 to 4 carbon atoms.
4. A mixture according to any one of Claims 1 to 3, containing at least one compound of formula Ia

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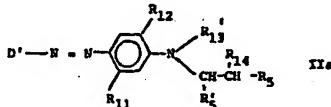


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30 and at least one compound of formula IIa

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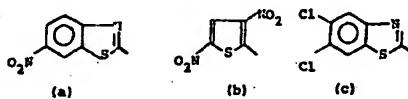
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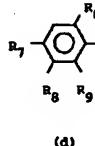
in which D' is a radical of formula (a), (b), (c) or (d)

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50 wherein

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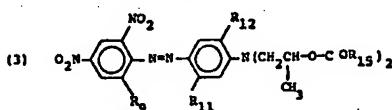
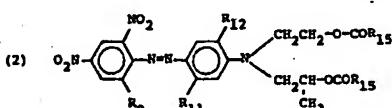
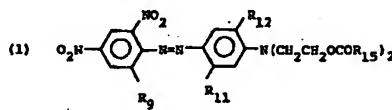
- R₆ is hydrogen, chlorine, bromine, iodine, nitro or cyano,
R₇ is nitro (C₁₋₁₂)alkylsulphonyl, aminosulphonyl, mono- or di(C₁₋₂)alkylaminosulphonyl or phenylazo in which the benzene ring is optionally substituted by up to three substituents selected from methyl, methoxy, chlorine, bromine and nitro,
55 R₈ is hydrogen, chlorine or bromine,
R₉ is hydrogen, chlorine, bromine, iodine, cyano or (C₁₋₂) alkylsulphonyl, with the proviso that one of R₈ and R₉ is hydrogen,
R₁₁ is hydrogen, methyl, methoxy, ethoxy, formylamino, alkyl (C₁₋₄)carbonylamino, alkyl (C₁₋₂)carbonylamino monosubstituted by hydroxyl, chlorine, bromine, cyano, phenyl, (C₁₋₃)alkyl xy or phen xy, alkoxy-(C₁₋₄)carbonylamino, th xycarbonylamino m n substituted by hydroxyl, (C₁₋₃)alkyl xy or (C₁₋₂) alkoxyeth xy, benz ylamino r (C₁₋₂)alkylsulphonylamino,
60 R₁₂ is hydrogen, methyl (C₁₋₂)alkoxy, (C₁₋₂)alk xyethoxy or (C₁₋₂) alkoxy monosubstituted by chlorine, bromine, cyano r phenyl,
R₁₃ is hydrogen, linear (C₁₋₄)alkyl, (C₂₋₃)-alkenyl, linear (C₂₋₃)alkyl mono-substituted by (C₁₋₂)alkoxy, cyano, formyloxy, alkyl (C₁₋₄) carbonyloxy, alk xy(C₁₋₃)carbonyl, benz yloxy or alkoxy (C₁₋₃)carbonyloxy,
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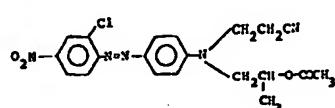
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- R₁₃ has one of the significances of R₁₃ but where the alkyl and substituted alkyl radicals are linear or branched,
- R₁₄ is formyloxy, alkyl (C₁₋₄)carbonyloxy, benzoxyloxy, phenoxyacetoxyl, alkoxy (C₁₋₄)carbonyloxy, alkoxy(C₁₋₄)carbonyl, alkoxy (C₁₋₂)ethoxycarbonyl, alkoxy (C₁₋₂)ethoxycarbonyloxy, alkoxy-(C₁₋₂)alkyl(C₁₋₂)carbonyloxy, mono- or di (C₁₋₄alkyl) aminocarbonyloxy or phenylaminocarbonyloxy wherein the benzene nucleus is optionally substituted by up to two substituents selected from chlorine, bromine, methyl and methoxy, whereby in such mixture D', R₁₁, R₁₂ and R₁₄ in the compound of formula Ia is identical with that of the compound of formula IIa.
- 5 10 5. A mixture according to Claim 4, in which D' is a group of formula (d).
6. A mixture according to Claim 4 or CLAIM %, IN WHICH R₆ is R₆', where R₆ is hydrogen, chlorine, bromine or nitro, R₇ is nitro, R₈ is hydrogen and R₉ is R₉' where R₉' is hydrogen, chlorine, bromine, cyano or methylsulphonyl.
7. A mixture according to any one of Claims 4 to 6, in which R₁₁ is R₁₁', where R₁₁' is hydrogen, methyl, 15 alkyl (C₁₋₂) carbonylamino, chloro- or bromoalkyl-(C₁₋₂) carbonylamino, alkoxy (C₁₋₂) carbonylamino or alkoxy-(C₁₋₂) ethoxycarbonylamino, R₁₂ is R₁₂', where R₁₂' is hydrogen, methoxy or ethoxy, R₁₃ is R_{13a}, wherein R_{13a} is linear (C₂₋₃) alkyl, allyl or linear (C₂₋₃) alkyl monosubstituted by cyano, alkyl (C₁₋₂) carbonyloxy, alkoxy (C₁₋₂)carbonyl or alkoxy (C₁₋₂) carbonyloxy, R₁₄ has one of the significances of R_{13a} but where the optionally substituted (C₂₋₃) alkyl may be linear or branched and R₁₄ is alkyl-(C₁₋₂) carbonyloxy or 20 alkoxy (C₁₋₂) carbonyloxy.
8. A mixture according to any one of Claims 1 to 6, in which R₆ is hydrogen and R₆' is methyl, R₁ is R₁₁, R₂ is R₁₂, R₃ is R₁₃, R₃' is R₁₃ all as defined in Claim 4 and R₄ is acyloxy.
9. A mixture according to Claim 8, in which R₄ is alkyl (C₁₋₂) carbonyloxy or alkoxy (C₁₋₂) carbonyloxy.
10. A mixture according to Claim 8 or Claim 9 in which R₁₁, R₁₂, R₁₃ and R₁₃' are as defined in Claim 7.
- 25 11. A mixture according to Claim 4, where D' is a radical (d) wherein R₆ is chlorine or bromine, R₇ is nitro, R₈, R₉, R₁₁ and R₁₂ are hydrogen, R₁₃ and R₁₃' are cyanoethyl, R₁₄ is alkyl (C₁₋₂) carbonyloxy, R₆ is methyl and R₆' is hydrogen;
12. A mixture according to Claim 4, where D' is a radical (d) wherein R₆, R₈, R₁₁ and R₁₂ are hydrogen, R₇ is nitro, R₉ is cyano, R₁₃ and R₁₃' are cyanoethyl, R₁₄ is alkyl (C₁₋₂) carbonyloxy, R₆ is methyl and R₆' is 30 hydrogen;
13. A mixture according to Claim 4, wherein D' is a radical (d) wherein R₆ and R₉, independently, are chlorine, or bromine,
- R₇ is nitro, R₈, R₁₁ and R₁₂ are hydrogen, R₁₃ and R₁₃' are cyanoethyl, R₁₄ is alkyl (C₁₋₂) - carbonyloxy, R₆ is methyl and R₆' is hydrogen;
- 35 14. A mixture according to Claim 4, comprising the following three dyestuffs

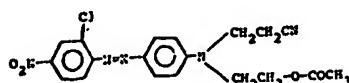


in which each R₉ is chlorine, bromine or iodine, especially bromine, each R₁₁ is alkyl (C₁₋₂)carbonylamino, each R₁₂ is (C₁₋₂) alkoxy, and each R₁₅ is (C₁₋₂) alkyl.

15. A mixture containing 15 to 85% dyestuff of formula

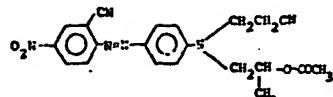


60 and 85% to 15% dyestuff of formula



16. A mixture containing 15% to 85% dyestuff of formula

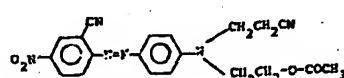
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and 85 to 15% dyestuff of formula

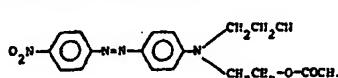
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17. A mixture containing 15 to 85% dyestuff of formula

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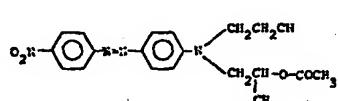


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and 85 to 15% dyestuff of formula

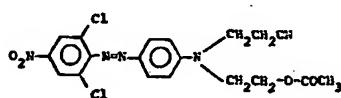
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18. A mixture containing 15 to 85% dyestuff of formula

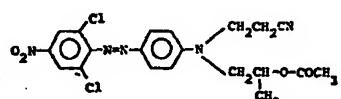
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and 85 to 15% dyestuff of formula

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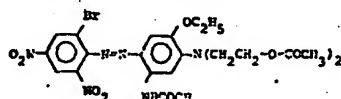


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19. A mixture containing approximately 25% dyestuff of formula

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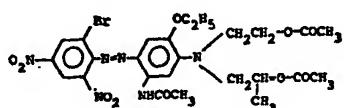


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approximately 50% dyestuff of formula

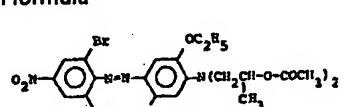
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and approximately 25% dyestuff of formula

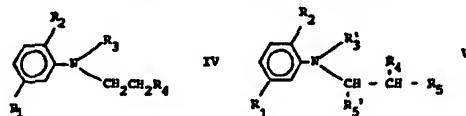
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20. A process for the preparation of a mixture according to Claim 1, comprising, coupling a diazotized amine with at least one compound of formula IV and at least one compound of formula V,

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whereby at least 15 mol % of a compound of formula IV and at least 15 mol % of a compound of formula V is present.

21. A process for the preparation of a mixture according to Claim 1, substantially as hereinbefore described with reference to any one of Examples 1 to 57.
- 5 22. A mixture of disperse dyestuffs whenever obtained by a process according to Claim 20 or Claim 21. 5
23. A process for dyeing or printing synthetic or semi-synthetic, hydrophobic, high-molecular weight organic textile substrates from aqueous suspension comprising employing a mixture according to any one of Claim 1 to 19 or 22 as dyeing agent.
- 10 24. A process for dyeing or printing synthetic or semi-synthetic, hydrophobic, high-molecular weight organic textile substrates substantially as hereinbefore described with reference to any one of Examples A to C. 10
25. Dyed or printed substrates whenever obtained by a process according to Claim 23 or 24.

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